



AP-2 α / β Monoclonal Antibody

Catalog No	YP-mAb-01531
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	TFAP2A/TFAP2B
Immunogen	The antiserum was produced against synthesized peptide derived from human AP-2. AA range:388-437
Specificity	AP-2 α / β Monoclonal Antibody detects endogenous levels of AP-2 α / β protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	$\geq 90\%$
Storage Stability	-20°C/1 year
Synonyms	Activating enhancer binding protein 2 alpha;Activating enhancer-binding protein 2-alpha;Activator protein 2;AP 2 transcription factor;AP 2alpha;AP-2;AP-2 transcription factor;AP2;AP2 Transcription Factor;AP2-alpha;AP2A_HUMAN;AP2TF;BOFS;FLJ51761;TFAP 2;TFAP 2A;TFAP2;TFAP2A;Transcription factor AP 2 alpha (activating enhancer binding protein 2 alpha);Transcription factor AP-2-alpha;Transcription factor AP2 alpha;Activating enhancer binding protein 2 beta;Activating enhancer-binding protein 2-beta;AP 2B;AP2 B;AP2-beta;AP2B;AP2B_HUMAN;AP2beta;MGC21381;OTTHUMP00000039925;PDA2;TFAP 2B;Tfap2b;Transcription factor AP 2 beta;Transcription factor AP-2-beta;Transcription factor AP2 beta;AP-2 α / β ;TFAP2A/TFAP2B;Transcription factor AP-2-alpha/beta;Activating enhancer-bind
Observed Band	49kD
Calculated Molecular Weight	49kD
Cell Pathway	Nucleus .
Tissue Specificity	Prostate,Teratocarcinoma,



Function

alternative products:Experimental confirmation may be lacking for some isoforms,disease:Defects in TFAP2A are the cause of branchiooculofacial syndrome (BOFS) [MIM:113620]; also known as branchial clefts with characteristic facies, growth retardation, imperforate nasolacrimal duct, and premature aging or lip pseudocleft-hemangiomas branchial cyst syndrome. BOFS is a rare autosomal dominant cleft palate craniofacial disorder with variable expressivity. The major features include cutaneous anomalies, ocular anomalies, characteristic facial appearance (malformed pinnae, oral clefts), and, less commonly, renal and ectodermal (dental and hair) anomalies.,domain:The WW-binding motif mediates interaction with WWOX.,function:Sequence-specific DNA-binding protein that interacts with inducible viral and cellular enhancer elements to regulate transcription of selected genes. AP-2 factors bind to

Background

transcription factor AP-2 alpha(TFAP2A) Homo sapiens The protein encoded by this gene is a transcription factor that binds the consensus sequence 5'-GCCNNNGGC-3'. The encoded protein functions as either a homodimer or as a heterodimer with similar family members. This protein activates the transcription of some genes while inhibiting the transcription of others. Defects in this gene are a cause of branchiooculofacial syndrome (BOFS). Three transcript variants encoding different isoforms have been found for this gene.[provided by RefSeq, Dec 2009],

matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis of various cells using AP-2α /β Monoclonal Antibody